

## CLAIMS

1. A method of treating a mammal for atherosclerotic disease comprising administering to the mammal a Factor XIIIa inhibitor.
2. A method according to claim 1, wherein the Factor XIIIa inhibitor is a Factor XIII(a) Lp(a)-matrix specific inhibitor.
3. A method according to claim 1, wherein the Factor XIIIa inhibitor is a Factor XIII(a) Lp(a)-fibrin specific inhibitor.
4. A method according to claim 1, wherein the mammal is a human.
5. A method of identifying an inhibitor of Factor XIIIa comprising:
  - (a) incubating an Lp(a) component, a matrix component, and Factor XIIIa in the presence or absence of a test inhibitor;
  - (b) determining whether complex formation between the Lp(a) component and the matrix component was inhibited in the presence of the test inhibitor; and
  - (c) identifying as a Factor XIIIa inhibitor the test inhibitor that inhibited complex formation.
6. A method according to claim 5, wherein the matrix component is selected from the group consisting of fibrin and a fibrin component.
7. A method of identifying a Factor XIIIa inhibitor comprising:
  - (a) incubating Factor XIIIa and a first substrate pair comprising an Lp(a) component and a matrix component in the presence or absence of a test inhibitor;
  - (b) incubating Factor XIIIa and a second substrate pair in the presence or absence of the test inhibitor, wherein the second substrate pair comprises any two components that are Factor XIIIa substrates for complex formation,
  - (c) determining whether inhibition of complex formation between the first substrate pair was greater than inhibition of complex formation between the second substrate pair; and
  - (d) identifying as a Factor XIIIa inhibitor the test inhibitor that provided greater inhibition of complex formation between the first substrate pair than between the second substrate pair.
8. A method according to claim 7, wherein the matrix component is selected from the group consisting of fibrin and a fibrin component.

9. A method according to claim 7, wherein the second substrate pair comprises a first member selected from the group consisting of fibrin and a fibrin component and a second member selected from the group consisting of fibrin and a fibrin component.

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